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NOTICE OF ALLOWANCE AND FEE(S) DUE

23413

09/18/2009

CANTOR COLBURN, LLP 20 Church Street 22nd Floor Hartford, CT 06103

EXAMINER WEATHERBY, ELLSWORTH ART UNIT PAPER NUMBER

3768 DATE MAILED: 09/18/2009

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/708,564	03/11/2004	Darin R. Okerlund	144726	2563

TITLE OF INVENTION: CARDIAC IMAGING SYSTEM AND METHOD FOR PLANNING MINIMALLY INVASIVE DIRECT CORONARY ARTERY

BYPASS SURGERY

APPLN. TYPE	SMALL ENTITY	ISSUE FEE DUE	PUBLICATION FEE DUE	PREV. PAID ISSUE FEE	TOTAL FEE(S) DUE	DATE DUE
nonprovisional	NO	\$1510	\$300	\$0	\$1810	12/18/2009

THE APPLICATION IDENTIFIED ABOVE HAS BEEN EXAMINED AND IS ALLOWED FOR ISSUANCE AS A PATENT. <u>PROSECUTION ON THE MERITS IS CLOSED</u>. THIS NOTICE OF ALLOWANCE IS NOT A GRANT OF PATENT RIGHTS. THIS APPLICATION IS SUBJECT TO WITHDRAWAL FROM ISSUE AT THE INITIATIVE OF THE OFFICE OR UPON PETITION BY THE APPLICANT. SEE 37 CFR 1.313 AND MPEP 1308.

THE ISSUE FEE AND PUBLICATION FEE (IF REQUIRED) MUST BE PAID WITHIN THREE MONTHS FROM THE MAILING DATE OF THIS NOTICE OR THIS APPLICATION SHALL BE REGARDED AS ABANDONED. THIS STATUTORY PERIOD CANNOT BE EXTENDED. SEE 35 U.S.C. 151. THE ISSUE FEE DUE INDICATED ABOVE DOES NOT REFLECT A CREDIT FOR ANY PREVIOUSLY PAID ISSUE FEE IN THIS APPLICATION. IF AN ISSUE FEE HAS PREVIOUSLY BEEN PAID IN THIS APPLICATION (AS SHOWN ABOVE), THE RETURN OF PART B OF THIS FORM WILL BE CONSIDERED A REQUEST TO REAPPLY THE PREVIOUSLY PAID ISSUE FEE TOWARD THE ISSUE FEE NOW DUE.

HOW TO REPLY TO THIS NOTICE:

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B. If the status above is to be removed, check box 5b on Part B -Fee(s) Transmittal and pay the PUBLICATION FEE (if required) and twice the amount of the ISSUE FEE shown above, or

If the SMALL ENTITY is shown as NO:

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II. PART B - FEE(S) TRANSMITTAL, or its equivalent, must be completed and returned to the United States Patent and Trademark Office (USPTO) with your ISSUE FEE and PUBLICATION FEE (if required). If you are charging the fee(s) to your deposit account, section "4b" of Part B - Fee(s) Transmittal should be completed and an extra copy of the form should be submitted. If an equivalent of Part B is filed, a request to reapply a previously paid issue fee must be clearly made, and delays in processing may occur due to the difficulty in recognizing the paper as an equivalent of Part B.

III. All communications regarding this application must give the application number. Please direct all communications prior to issuance to Mail Stop ISSUE FEE unless advised to the contrary.

IMPORTANT REMINDER: Utility patents issuing on applications filed on or after Dec. 12, 1980 may require payment of maintenance fees. It is patentee's responsibility to ensure timely payment of maintenance fees when due.

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Complete and send this form, together with applicable fee(s), to: Mail Mail Stop ISSUE FEE

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Hartford, CT 06	103							(Depositor's name)
								(Signature)
								(Date)
APPLICATION NO.	FILING DATE		FIRST NAMED INVEN	TOR		ATTO:	RNEY DOCKET NO.	CONFIRMATION NO.
10/708,564	03/11/2004		Darin R. Okerlund	d	•		144726	2563
ITLE OF INVENTION YPASS SURGERY	: CARDIAC IMAGING	SYSTEM AND METHO	DD FOR PLANNING	MIN	IMALLY INVASI	VE DI	RECT CORONARY A	ARTERY
APPLN. TYPE	SMALL ENTITY	ISSUE FEE DUE	PUBLICATION FEE D	UE	PREV. PAID ISSUE	FEE	TOTAL FEE(S) DUE	DATE DUE
nonprovisional	NO	\$1510	\$300		\$0		\$1810	12/18/2009
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WEATHERBY,	ELLSWORTH	3768	600-425000					
Change of corresponde FR 1.363). Change of corresp Address form PTO/SI "Fee Address" ind PTO/SB/47; Rev 03-0 Number is required.	(1) the names of u or agents OR, alter (2) the name of a s registered attorney	of a single firm (having as a member a orney or agent) and the names of up to patent attorneys or agents. If no name is						
PLEASE NOTE: Unl recordation as set fort (A) NAME OF ASSIG	less an assignee is ident h in 37 CFR 3.11. Comp GNEE		data will appear on the Tasubstitute for filing (B) RESIDENCE: (C)	he pa g an a	tent. If an assigne ssignment. and STATE OR Co	TNUC	RY)	ocument has been filed for
lease check the appropr	rate assignee category or	categories (will not be pr	inted on the patent):		Individual 🖵 Co.	rporati	on or other private gro	up entity Government
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	tus (from status indicated s SMALL ENTITY statu	,	☐ b. Applicant is no	long	er claiming SMAL	L ENT	ΓΙΤΥ status. See 37 CF	FR 1.27(g)(2).
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APPLICATION NO.	ION NO. FILING DATE FIRST NAMED INVENTOR		ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/708,564	03/11/2004	Darin R. Okerlund	144726	2563
23413 75	590 09/18/2009		EXAM	INER
CANTOR COLE	BURN, LLP	WEATHERBY,	ELLSWORTH	
20 Church Street			ART UNIT	PAPER NUMBER
22nd Floor Hartford, CT 0610	3		3768 DATE MAILED: 09/18/200	9

Determination of Patent Term Adjustment under 35 U.S.C. 154 (b)

(application filed on or after May 29, 2000)

The Patent Term Adjustment to date is 791 day(s). If the issue fee is paid on the date that is three months after the mailing date of this notice and the patent issues on the Tuesday before the date that is 28 weeks (six and a half months) after the mailing date of this notice, the Patent Term Adjustment will be 791 day(s).

If a Continued Prosecution Application (CPA) was filed in the above-identified application, the filing date that determines Patent Term Adjustment is the filing date of the most recent CPA.

Applicant will be able to obtain more detailed information by accessing the Patent Application Information Retrieval (PAIR) WEB site (http://pair.uspto.gov).

Any questions regarding the Patent Term Extension or Adjustment determination should be directed to the Office of Patent Legal Administration at (571)-272-7702. Questions relating to issue and publication fee payments should be directed to the Customer Service Center of the Office of Patent Publication at 1-(888)-786-0101 or (571)-272-4200.

	Application No.	Applicant(s)					
	10/708,564	OKERLUND ET AL.					
Notice of Allowability	Examiner	Art Unit					
	ELLSWORTH WEATHERBY	3768					
The MAILING DATE of this communication appear All claims being allowable, PROSECUTION ON THE MERITS IS herewith (or previously mailed), a Notice of Allowance (PTOL-85) NOTICE OF ALLOWABILITY IS NOT A GRANT OF PATENT RIOF of the Office or upon petition by the applicant. See 37 CFR 1.313	(OR REMAINS) CLOSED in this ap or other appropriate communicatio IGHTS. This application is subject	oplication. If not included n will be mailed in due course. THIS					
1. This communication is responsive to <u>11/25/2008</u> .							
2. X The allowed claim(s) is/are <u>Claims 1-4, 6, 8-13, 16-20, 22-</u>	<u>26, and 28-30</u> .						
 3. ☐ Acknowledgment is made of a claim for foreign priority ur a) ☐ All b) ☐ Some* c) ☐ None of the: 1. ☐ Certified copies of the priority documents have 							
□ Certified copies of the priority documents have □ Certified copies of the priority documents have							
Copies of the certified copies of the priority do	• • • • • • • • • • • • • • • • • • • •						
International Bureau (PCT Rule 17.2(a)).	odinente nave been received in tine	Thational stage application from the					
* Certified copies not received:							
Applicant has THREE MONTHS FROM THE "MAILING DATE" of this communication to file a reply complying with the requirements noted below. Failure to timely comply will result in ABANDONMENT of this application. THIS THREE-MONTH PERIOD IS NOT EXTENDABLE.							
4. A SUBSTITUTE OATH OR DECLARATION must be submitted. Note the attached EXAMINER'S AMENDMENT or NOTICE OF INFORMAL PATENT APPLICATION (PTO-152) which gives reason(s) why the oath or declaration is deficient.							
5. CORRECTED DRAWINGS (as "replacement sheets") mus	st be submitted.						
(a) ☐ including changes required by the Notice of Draftsperson's Patent Drawing Review (PTO-948) attached							
1) 🔲 hereto or 2) 🔲 to Paper No./Mail Date							
(b) ☐ including changes required by the attached Examiner's Paper No./Mail Date	(b) ☐ including changes required by the attached Examiner's Amendment / Comment or in the Office action of						
Identifying indicia such as the application number (see 37 CFR 1 each sheet. Replacement sheet(s) should be labeled as such in t							
6. DEPOSIT OF and/or INFORMATION about the deposit of BIOLOGICAL MATERIAL must be submitted. Note the attached Examiner's comment regarding REQUIREMENT FOR THE DEPOSIT OF BIOLOGICAL MATERIAL.							
Attachment(s) 1. ☐ Notice of References Cited (PTO-892)	5. ☐ Notice of Informal I	Datant Application					
 Induce of References Cited (PTO-092) Induce of References Cited (PTO-	6. ☐ Interview Summary	• •					
	Paper No./Mail Da	ate					
3. 🛮 Information Disclosure Statements (PTO/SB/08), Paper No./Mail Date <u>See Continuation Sheet</u>	7. ⊠ Examiner's Amend	ment/Comment					
Examiner's Comment Regarding Requirement for Deposit of Biological Material	8. 🛛 Examiner's Statem	ent of Reasons for Allowance					
	9.						

Continuation of Attachment(s) 3. Information Disclosure Statements (PTO/SB/08), Paper No./Mail Date: 12/28/2006; 10/26/2006; 8/16/2006; 3/17/2006; 12/20/2004; 9/15/2004; 3/11/2004.

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EXAMINER'S AMENDMENT

1. An examiner's amendment to the record appears below. Should the changes and/or additions be unacceptable to applicant, an amendment may be filed as provided by 37 CFR 1.312. To ensure consideration of such an amendment, it MUST be submitted no later than the payment of the issue fee.

Authorization for this examiner's amendment was given in a telephone interview with John Buckert (Reg. 44,572) on 6/5/2009.

The application has been amended as follows:

Claim 1. (Currently Amended): A method for planning <u>a</u> minimally invasive direct coronary artery bypass (MIDCAB) for a patient, the method comprising:

obtaining acquisition data from a medical imaging system;

generating a 3D model of coronary arteries <u>and one or more cardiac chambers of</u> interest of the patient from said acquisition data, prior to performing said MIDCAB;

identifying one or more anatomical landmarks on said 3D model and inserting corresponding translucent geometric markers thereat, utilizing user input at an operator console:

saving views of said 3D model in a database;

measuring sizes of lesions and a number of the lesions in the coronary arteries utilizing said 3D model;

registering said saved views of said 3D model on a workstation of an interventional system, said saved views of said 3D model having said translucent geometric markers; and, said registering including importing said saved views of said 3D model having said translucent geometric markers to the coordinate system of said interventional system using said one or more anatomical landmarks;

visualizing one or more of said registered saved views on a display screen of said interventional system; and

utilizing the interventional system to quantify distance and location information for a cardiac point of interest prior performing said MIDCAB; and

identifying an incision location and path for said MIDCAB based on said quantified distance and location information for said cardiac point of interest.

Claim 9. (Currently Amended): A method for planning <u>a</u> minimally invasive direct coronary artery bypass (MIDCAB) for a patient, the method comprising:

obtaining acquisition data from a medical imaging system using a protocol directed toward the coronary arteries and left ventricle;

segmenting said acquisition data using a 3D protocol so as to visualize <u>the</u> <u>coronary arteries and the left ventricle;</u>

generating a 3D model of the coronary arteries and the left ventricle of the patient from said acquisition data, prior to performing said MIDCAB;

identifying one or more anatomical landmarks on said 3D model and inserting corresponding translucent geometric markers thereat, utilizing user input at an operator console;

saving views of said 3D model in a database;

measuring sizes of lesions and a number of lesions in the coronary arteries utilizing said 3D model and identifying, from said 3D model, an orientation and any anomalies associated with the coronary arteries;

registering said saved views of said 3D model on a workstation of an interventional system, said saved views of said 3D model having said translucent geometric markers, said registering including, transforming said saved views of said 3D model having said translucent geometric markers to the coordinate system of said interventional system using said one or more anatomical landmarks;

visualizing one or more of said registered saved views on a display screen of said interventional system; and

identifying, from said 3D model, orientation and any anomalies associated with the coronary arteries and the left ventricle;

using said identified orientation and anomalies associated with the coronary arteries and said registered saved views to determine appropriate sites for incisions for targeted MIDCAB.

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Claim 16. (Currently Amended): A method for planning <u>a</u> minimally invasive direct coronary artery bypass (MIDCAB) for a patient, the method comprising:

obtaining acquisition data from a cardiac computed tomography (CT) imaging system using a protocol directed toward coronary arteries and one or more cardiac chambers of interest;

segmenting said acquisition data using a 3D protocol so as to visualize the coronary arteries, including interior views of the coronary arteries;

generating a 3D model of the coronary arteries of the patient <u>from said</u> acquisition data, prior to performing said MIDCAB;

identifying one or more anatomical landmarks on said 3D model and inserting corresponding translucent geometric markers thereat, utilizing user input at an operator console;

saving views of said 3D model in a database;

measuring sizes of lesions and a number of the lesions in the coronary arteries utilizing said 3D model and identifying, from said 3D model, an orientation and any anomalies associated with the coronary arteries;

registering said saved views of said 3D model on <u>a</u> fluoroscopy system, said saved views of said 3D model having said geometric markers, <u>said registering including transforming said saved views of said 3D model having said translucent geometric markers to the coordinate system of the fluoroscopy system using said one or more anatomical landmarks; and</u>

visualizing one or more of said registered saved views with said fluoroscopy system; and

<u>quantifying distance and location information for a cardiac point of interest; and identifying an incision location and path for said MIDCAB based on said quantified distance and location information for the cardiac point of interest.</u>

Claim 22. (Currently Amended): A system for planning <u>a</u> minimally invasive direct coronary artery bypass (MIDCAB) for a patient, comprising:

a medical imaging system for generating acquisition data;

an image generation subsystem for receiving said acquisition data and generating one or more images and a 3D model of coronary arteries and one or more cardiac chambers of interest of the patient, the image generation subsystem further configured to automatically measuring measure sizes of lesions and a number of the lesions in the coronary arteries utilizing said 3D model;

an operator console <u>configured to receive</u> for receiving user input to identify one or more anatomical landmarks on said one or more images or said 3D model and to insert corresponding geometric markers thereat, said console further configured to save <u>views of said 3D model having said geometric markers to a database</u>;

a workstation of an interventional system configured to receive said saved views
of said 3D model having said geometric markers from said database, where said
workstation including includes post processing software stored on a computer readable
medium for registering said images of said saved views of said 3D model on an

interventional system , said 3D model having said geometric markers; and by transforming said saved views of said 3D model having said geometric markers to the coordinate system of an interventional system using said one or more anatomical landmarks;

wherein said workstation is configured for visualizing one or more of said registered images therewith, quantifying distance land location information for a cardiac point of interest, and identifying an incision location and path for MIDCAB based on said quantified distance and location information for said cardiac point of interest.

wherein said workstation of said interventional system is configured to:

import said registered saved views of said 3D model having said geometric

markers;

visualize said registered saved views of said 3D model having said geometric markers; and

utilize said registered saved views of said 3D model having said geometric markers to quantify distance and location information for a cardiac point of interest to identify an incision location and path for MIDCAB.

Allowable Subject Matter

The following is an examiner's statement of reasons for allowance: The prior art of record does not disclose or suggest, *inter alia*, a method for planning minimally invasive direct coronary artery bypass (MIDCAB) for a patient, the method comprising: obtaining acquisition data from a medical imaging system; generating a 3D model of the

left ventricle and thoracic wall of the patient from the acquisition data, prior to performing a MIDCAB procedure on the patient; identifying one or more left ventricle anatomical landmarks on said 3D model and inserting geometric markers therein corresponding to selected ones of said anatomical landmarks; registering saved views of said 3D model on an interventional system; and visualizing one or more of said registered saved views with said interventional system; and identifying sizes and numbers of lesions utilizing the 3D model.

Concerning the section 102(b) rejection using Vesely, the examiner agrees with applicant in that the cited reference does not teach or suggest a method or system for planning MIDCAB. Here, the steps of: identifying one or more left ventricle anatomical landmarks on said 3D model and inserting geometric markers therein corresponding to selected ones of said anatomical landmarks; registering saved views of said 3D model on an interventional system; and identifying sizes and numbers of lesions utilizing the 3D model are not taught or suggested by the prior art. Based on the above observations, claims 1-4, 6, 8-13, 16-20, 22-26, and 28-30 are allowable.

Any comments considered necessary by applicant must be submitted no later than the payment of the issue fee and, to avoid processing delays, should preferably accompany the issue fee. Such submissions should be clearly labeled "Comments on Statement of Reasons for Allowance."

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Terminal Disclaimer

2. The terminal disclaimer filed on 11/25/2008 disclaiming the terminal portion of any patent granted on this application which would extend beyond the expiration date of USPN 7,343,196 has been reviewed and is accepted. The terminal disclaimer has been recorded.

3. The terminal disclaimer filed on 11/25/2008 disclaiming the terminal portion of any patent granted on this application which would extend beyond the expiration date of 7,346,381 has been reviewed and is accepted. The terminal disclaimer has been recorded.

Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to ELLSWORTH WEATHERBY whose telephone number is (571) 272-2248. The examiner can normally be reached on M-F 8:30 a.m. - 5:00 p.m..

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Long Le can be reached on (571) 272-0823. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/EW/

/Long V Le/ Supervisory Patent Examiner, Art Unit 3768